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CAPILLARY REACTOR DISTRIBUTION DEVICE AND METHOD

Abstract:

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A capillary reactor distribution device comprising first and second capillary pathways (2, 3) which meet at a junction (5) and a third capillary pathway (4) which leads away from the junction (5), the capillary pathways (2, 3, 4) being dimensioned such that, when first and second immiscible fluids (14, 15) are fed along respectively the first and second capillary pathways (2, 3) under predetermined laminar flow conditions, the first and second fluids (14, 15) chop each other into discrete slugs (16, 17) which pass along the third capillary pathway (4). Molecular mixing between the fluids (14, 15) takes place by way of axial diffusion between adjacent slugs (16, 17) and by way of internal circulation within each slug (16, 17) as the slugs (16, 17) progress along the third capillary pathway (4).

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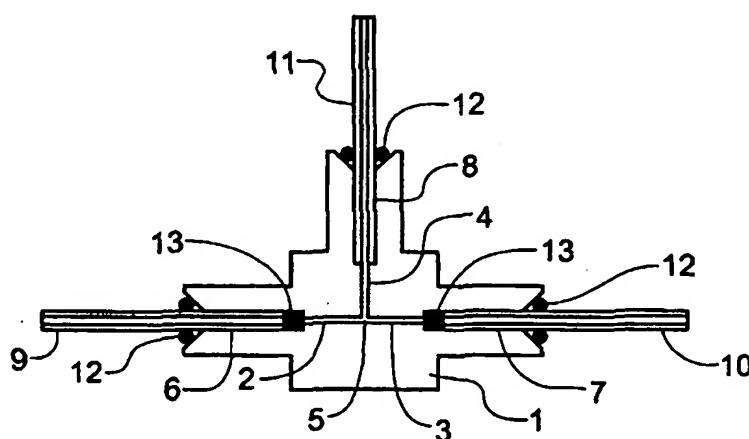
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(54) Title: CAPILLARY REACTOR DISTRIBUTION DEVICE AND METHOD



(57) Abstract: A capillary reactor distribution device comprising first and second capillary pathways (2, 3) which meet at a junction (5) and a third capillary pathway (4) which leads away from the junction (5), the capillary pathways (2, 3, 4) being dimensioned such that, when first and second immiscible fluids (14, 15) are fed along respectively the first and second capillary pathways (2, 3) under predetermined laminar flow conditions, the first and second fluids (14, 15) chop each other into discrete slugs (16, 17) which pass along the third capillary pathway (4). Molecular mixing between the fluids (14, 15) takes place by way of axial diffusion between adjacent slugs (16, 17) and by way of internal circulation within each slug (16, 17) as the slugs (16, 17) progress along the third capillary pathway (4).

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CAPILLARY REACTOR DISTRIBUTION DEVICE AND METHOD

The present invention relates to a capillary reactor distribution device and method and in particular to a distribution device and method for interfacing at least two
5 immiscible fluids.

Improved methods of manufacturing at microscales have opened up new opportunities for the development of compact, efficient and highly controllable reactors. Rapid mass and heat transfer between fluids may now be engineered into
10 reactors by the use of small dimensions for fluid transport. Short path lengths for thermal and molecular diffusion can provide an ideal environment for rapid exothermic/endothermic reactions while maintaining a laminar flow regime. Scale requirements for efficient mixing by diffusion can be calculated using the following equation (Crank, J., 1975, "The Mathematics of Diffusion", 2nd edition, Clarendon
15 Press, Oxford):

$$d = (Fo \cdot Dt)^{0.5} \quad (1)$$

where D is the diffusivity of the reacting molecules in the fluid, t is the residence
20 time and Fo the Fourier number determining the level of mixing. For most systems, Fo = 1 would be chosen. Reactions that may benefit most from this technique are those where phases cannot mix to form a single phase, such as liquid-gas, liquid-solid, gas-solid and immiscible liquid flow.

25 Two general methods are available for efficiently contacting two immiscible liquid streams within a microreactor. The first is the use of parallel liquid streams as described in WO 97/39814 and WO 99/22858 where diffusion is perpendicular to the flow direction. The second is the use of dispersed/continuous phase flow where one
30 phase is in the form of small droplets within the other phase or slug flow where each phase is in the form of a series of slugs. Diffusive mass transfer may be aided by internal circulation within the droplets generated by the shear flow as demonstrated

by Clift, R., Grace, J.R. and Weber, M.E. (1978, "Bubbles, drops and particles", Academic Press, New York).

Several benefits and drawbacks accompany these two techniques. In the use of
5 parallel flow, it is difficult to achieve stability and similar residence times for liquids
of significantly different viscosity or flow rate. On the other hand, for droplet flow,
velocities of the dispersed and continuous phases remain similar, and a wider range
of flow rate ratios can be tolerated. However, parallel flow has the advantage of easy
10 bulk separation of the liquids after reaction, whereas droplets and slugs need to be
separated by way of centrifugal or gravitational action.

Typical diffusion rates within liquids are in the range $10^{-8} \text{ m}^2 \text{ s}^{-1}$ to $10^{-9} \text{ m}^2 \text{ s}^{-1}$, and
therefore, from equation (1) above, length scales for diffusion within the reactor are
required to be of the order of $100\mu\text{m}$ for rapid reactions requiring 1 to 10s residence
15 time. However, mass transfer enhancement due to internal vortices in droplet/slug
flow augment this process and allow larger channel dimensions to be used while
maintaining fast reaction times.

It is known from US 5,921,678 to provide a microfluidic sub-millisecond mixer in
20 which two capillary pathways meet head-on and in which a third capillary pathway
leads away from the junction of the first two, thereby forming a T-junction. Two
miscible fluids are then directed along the first two capillary pathways so as to meet
at the junction, mix in turbulent conditions, and then flow along the third capillary
pathway where reaction takes place. The third capillary pathway is very short, so as
25 to constrain reaction time to sub-millisecond timescales, before the reactants are
quenched and then separated. This prior art mixer is not suited for use with
immiscible fluids.

According to a first aspect of the present invention, there is provided a capillary
30 reactor distribution device comprising first and second capillary pathways which
meet at a junction and a third capillary pathway which leads away from the junction,

the capillary pathways being dimensioned such that, when first and second immiscible fluids are fed along respectively the first and second capillary pathways under predetermined laminar flow conditions, the first and second fluids chop each other into discrete slugs which pass along the third capillary pathway.

5

- According to a second aspect of the present invention, there is provided a method for contacting two immiscible fluids, wherein a first fluid is fed under laminar flow conditions along a first capillary pathway and a second fluid is fed under laminar flow conditions along a second capillary pathway, the first and second capillary pathways meeting at a junction having a third capillary pathway leading away therefrom, and wherein the flow conditions in each of the first and second capillary pathways are selected such that the first and second fluids chop each other into discrete slugs which pass along the third capillary pathway.
- 10
- 15 In general, the two immiscible fluids are both in the liquid phase, although liquid/gas, solid/liquid/liquid and solid/liquid/gas reactions may be performed in the device and method of the present invention. In the case of solid/liquid/liquid and solid/liquid/gas reactions, the solid phase may be coated on a surface of the third capillary pathway, for example in the form of a catalyst coating for liquid/liquid or liquid/gas reactions.
- 20

- In preferred embodiments, the first and second capillary pathways meet substantially head-on at the junction, although for some applications the first and second capillary pathways may be arranged to meet at angles other than substantially 180 degrees.
- 25 For example, the first and second capillary pathways may be arranged to meet at an angle of substantially 90 degrees, at an angle between 90 and 180 degrees, or at an angle between 0 and 90 degrees. In some embodiments, the first and second capillary pathways may be arranged to meet at an angle of 90 degrees to 300 degrees, and the third capillary pathway may lie substantially midway between the first and second capillary pathways or at substantially 180 degrees to the first capillary pathway.
- 30

Advantageously, the capillary pathways are formed in or lined with a non-stick or low surface energy material, such as a fluoropolymer (e.g. PTFE or PVDF).

- It is believed that the mechanism whereby the first fluid chops the second fluid into discrete slugs is as follows. Assuming that each fluid approaches the junction at a constant flow rate, and considering the case where the first and second capillary pathways meet substantially head-on, the first fluid flows preferentially into the third, exit capillary pathway while the second fluid forms an interface at the junction. The interface is moved into the third capillary pathway by the driving pressure from the first fluid supply aided by viscous shear from the second fluid. When the interface grows to a size that blocks the first fluid from entering the third pathway, the process switches and the second fluid flows preferentially into the third pathway while the first fluid forms an interface, which then moves into the third pathway before the process switches back again. This alternating flow of the first and second fluids generates a series of slugs in the third pathway. The lengths of the slugs is believed to be most significantly governed by the ratios of the widths of the inlet and outlet capillary pathways, the surface energy of the walls of the capillary pathways and the ratio of the first and second fluid flow rates. In particular, the ratio of slug lengths is generally substantially the same as the ratio of fluid flow rates. The lengths of the slugs are governed also to a lesser extent by the total fluid flow and the viscosities of the first and second fluids, and also by interfacial phenomena. Production of slugs is most preferably achieved in materials which do not have very low contact angles with either of the fluids.
- As the slugs progress along the third capillary pathway, mixing of the first and second fluids on a molecular level is achieved by both axial diffusion between adjoining slugs and also by internal circulation within each slug, the latter process generally being the dominant one. Both forms of mixing will generally increase as slug length is reduced. For rapid mixing, the smallest slug length should be of the order of the width of the third capillary pathway, and the longest slug length not greater than 100 times the width thereof, and preferably not greater than 10 times the

width thereof. It is particularly preferred that the longest slug length is not greater than twice the width of the third capillary pathway.

- 5 The device of the present invention may comprise a solid block of any appropriate material having the capillary pathways bored thereinto. Fluid may be pumped to the block and removed therefrom by way of standard capillary tubes which are connected to the capillary pathways bored into the block. The capillary tubes may connect to the capillary pathways at or near external surfaces of the block, or may connect thereto within the body of the block, near to the junction. Alternatively, passages 10 may be bored into the block so as snugly to receive the capillary tubes, the junction being defined by the ends of the capillary tubes themselves where they meet within the block. Preferably, O-ring or similar seals are provided where the capillary tubes enter the block so as to prevent pressure losses as fluid is pumped towards the 15 junction. Advantageously, the internal volume of the capillary pathways within the block is kept as small as possible.

Typical flow rates through the device of the present invention range from 10nls^{-1} to $100\mu\text{l s}^{-1}$, with preferred flow rates ranging from 100nls^{-1} to $10\mu\text{l s}^{-1}$. Flow rate ratios 20 between the fluids in the first and second capillary pathways are advantageously not greater than 10:1 and are preferably close to unity for high mixing efficiency.

Alternatively, the device of the present invention may be formed by at least two generally laminar plates mounted one directly on top of the other such that a surface 25 of one plate contacts a surface of the other plate, at least one of the surfaces being provided with features serving to define the capillary pathways. The plates will generally be in registration with each other. The at least one surface may include channels or ridge-like protrusions or both, such that when the surfaces of the plates are contacted, the required capillary pathways are defined between the plates. The 30 channels and/or the protrusions may be formed by an etching process, or may be micromachined or moulded. Further plates with suitable surface features may be

stacked on top of the at least two plates so as to produce a multi-layer device.

Either the first or the second capillary pathway or both may be provided with a fluid filter to help prevent stray particles from entering the device and which may block the 5 capillary pathways. In the solid block embodiment of the present invention, the fluid filter is preferably located between either one or both of the input capillary tubes and the first or second capillary pathways.

The device and method of the present invention is particularly useful for conducting 10 reactions between organic and aqueous liquids, for example the nitration of benzene and toluene as discussed hereinafter. Other applications include rapid mass transfer for liquid-liquid extraction and small volume reaction testing for analytical purposes.

For a better understanding of the present invention and to show how it may be carried 15 into effect, reference shall now be made by way of example to the accompanying drawings, in which:

FIGURES 1 to 3 show three alternative configurations of the distribution device of the present invention;

20 FIGURES 4 to 7 show a proposed mechanism for slug formation within the device of the present invention;

FIGURE 8 shows a series of liquid slugs within a capillary tube;

25 FIGURE 9 shows a capillary reactor including a device of the present invention;

FIGURE 10 is a graph of reaction rates against temperature for benzene nitration in the reactor of Figure 9;

30 FIGURE 11 is a graph of reaction rates for benzene nitration in the reactor of Figure

9 at different flow velocities;

FIGURE 12 is a graph of nitrotoluene concentration against acid/organic flow ratio for toluene nitration along different reaction pathway lengths in the reactor of Figure

5 9;

FIGURE 13 is a graph of transfer times for acetic acid from organic to aqueous phase in a 0.38mm x 0.38mm glass channel;

10 FIGURE 14 shows an alternative embodiment of the device of the present invention; and

FIGURE 15 shows the flow dynamics of slug propagation along a capillary pathway.

15 Figure 1 is a section through a PTFE block 1 into which first, second and third tubular capillary pathways 2, 3, 4 of diameter 0.5mm and length 5mm have been bored. The first and second capillary pathways 2, 3 meet head-on at a junction 5, and the third capillary pathway 4 leads away from the junction 5 substantially at right angles to the first and second capillary pathways 2, 3. Boreholes 6, 7, 8 are provided
20 so as to allow PTFE capillary tubes 9, 10, 11 with internal diameters of 0.15mm to be snugly inserted into the block 1 and to connect respectively to the capillary pathways 2, 3, 4. Each capillary tube 9, 10, 11 is provided with an O-ring seal 12 where it enters the block 1 so as to reduce pressure losses within the block 1, and filters 13 are provided where the feed capillary tubes 9, 10 connect with the first and second
25 capillary pathways 2, 3.

An alternative embodiment is shown in Figure 2, where boreholes 6, 7, 8 extend to a middle portion of the block 1 so as to allow the capillary tubes 9, 10, 11 with internal diameters of 0.15mm to meet and form the junction 5.

30

A further alternative embodiment is shown in Figure 3, where the capillary pathways

2, 3, 4 of diameter 0.8mm extend from the junction 5 to external surfaces of the block 1, where the capillary tubes 9, 10, 11 are connected.

It is to be noted that in some embodiments, the positions of capillary pathway 3 and 5 capillary tube 10 may be swapped with those of capillary pathway 4 and capillary tube 11, such that the two feed capillary pathways 2, 3 meet substantially at right angles.

By using a syringe driver (not shown) to inject dyed kerosene 14 along capillary tube 10 9 and thence capillary pathway 2, and water 15 along capillary tube 10 and thence capillary pathway 3, a series of slugs 16, 17 were formed in capillary pathway 4 and thence capillary tube 11, as shown in Figures 4 to 7 and 8.

The mechanism for slug 16, 17 formation is shown in Figures 4 to 7. In Figure 4, 15 water 15 flows preferentially from the capillary pathway 3 into the capillary pathway 4, while kerosene 14 forms an interface 18 at the junction 5. Due to the driving pressure behind the kerosene 14, the interface 18 is moved into the junction 5 and towards the capillary pathway 4, aided by viscous shear from the water 15 as shown in Figures 5 and 6. When the interface 18 has completely moved into the capillary 20 pathway 4, as shown in Figure 7, the flow of water 15 is blocked and kerosene 14 then flows preferentially into the capillary pathway 4, with the water 15 forming an interface 18'. The process is then reversed until the interface 18' moves into the capillary pathway 4 and water 15 again flows preferentially into the capillary pathway 4. The alternating movement of the interface 18, 18' causes a series of kerosene 25 slugs 16 and water slugs 17 to be formed in the capillary pathway 4 and thence the capillary tube 11 as shown in Figure 8.

Flow rates of 0.8 to 13 μ ls⁻¹ were tested, with aqueous/organic flow ratios of 2:1 and 1:1. The embodiment of Figure 1 was found to produce slug 16, 17 lengths of 2.1 to 30 5.5mm, that of Figure 2 to produce lengths of 0.3 to 0.9mm and that of Figure 3 to produce lengths of 18 to 30mm. This indicates that the low internal flow volume of

the Figure 2 embodiment helps to produce short slug 16, 17 lengths.

Figure 9 shows a reactor comprising an aqueous phase pump 19 and an organic phase pump 20, respectively connected to capillary tubes 9, 10 which then pass into a distribution device 1 of the type shown in Figure 2. An output capillary tube 11 passes from the device 1 and through a heater 21, inside which the capillary tube 11 is coiled for efficient use of space. The capillary tube 11 then passes from the heater 21 to a collection bottle 22. In the following examples, an aqueous phase reactant was pumped by pump 19 along capillary tube 9 and an organic phase reactant by pump 20 along capillary tube 10. Slugs (not shown) of aqueous phase and organic phase reactant were formed in the capillary tube 11 by the device 1, and then passed along the capillary tube 11, through the heater 21 and thence to the collection bottle 22 which contained solvents to halt the reaction between the reactants and to dilute the organic phase reactant. Analysis of the organic conversions discussed in the following examples was performed using gas chromatography.

Example 1: Benzene nitration

Distributors having capillary tubes made out of 316 stainless steel with respectively 127 μ m, 178 μ m and 254 μ m bore sizes were constructed. A syringe driver was used to supply the liquids for the reaction and a heating bath was used to control the reactor temperature. The nitration reaction involved contacting a stream of benzene with a stream of nitric and concentrated sulphuric acids. Various acid strengths and reactor temperatures were used in the nitration work and comparisons made of the reaction rate and by-product formation. A shell reaction model was used in calculating the reaction rate for the process. This assumes that nitration takes place in a acid boundary layer surrounding the organic phase. For this model mass transfer into the region and kinetic reaction rate within the region are equally important in the overall observed rate. The resulting equation governing this process can be written as a 1.5th order reaction, as shown in equation (2), where X is the proportion of the initial nitric acid remaining at time t.

$$\frac{dX}{dt} = -CX^{1.5}$$

(2)

The value of constant C is determined by the mass transfer rate into the reaction zone and kinetic reaction rate within the zone. Integration of equation (2) yields the following equation for nitric acid concentration at time t.

$$X = \left(1 + \frac{Ct}{2}\right)^{-2}$$

(3)

One method of characterising the nitration process is by comparison of the indicated initial reaction rates. This is defined as the reaction rate at the start of the process and can be calculated from equations (2) and (3) as,

$$InitialRate = \frac{dX}{dt} \Big|_{t=0} = C = \frac{2(X^{-1/2} - 1)}{t}$$

(4)

- 10 where X is the measured value at time t. A similar formula can be obtained for the organic reaction rate by substitution of X with the proportion of non-nitrated organic remaining. A comparison of the reaction rate observed for 127 μ m and 254 μ m bore capillary tubing under similar conditions is shown in Figure 10. A high sulphuric acid concentration was used to ensure fast nitration kinetics and promote a mass transfer limited regime. Comparing the results for the two capillary diameters clearly shows enhanced performance at the smaller scale implying improved mixing.
- 15

Figure 11 illustrates the influence of capillary flow rate on organic reaction rate. An enhancement in reaction rate was observed when faster flow was applied to the reactor especially for the conditions with the fastest kinetics. This would indicate that increased velocity was leading to increased mixing. The primary source of this improvement is most likely due to increased internal circulation within the liquids, although some variation in slug length may also be contributing.

Example 2: Toluene nitration

Recent work has examined the nitration of toluene using a PTFE capillary reactor.

- 5 The use of PTFE gave a more corrosion resistant system with less chance of blockage. Blockages were found to occur occasionally in the stainless steel system between runs probably due to sulphuric acid corrosion. However, no such problems occurred with the PTFE based system. Two HPLC pumps were used to supply the flow to the reactor with a greater run time capability.

10

- Toluene nitration was performed using 150 μm bore tubing using a range of acid strengths and reactor temperatures. Results showed a lower influence of temperature on reaction rate than benzene nitration when temperatures of greater than 75°C were used. Typical nitric acid reaction rates for the system are shown in Table 1 for a range of acid and organic flow ratios. Observed rates were generally higher than for benzene under similar conditions.

Acid:Organic Flow Ratio	Reaction Rate at 25°C (min ⁻¹)	Reaction Rate at 60°C (min ⁻¹)
2:1	3.27	6.10
3:1	2.81	6.12
5:1	2.25	4.39
7:1	1.65	4.17

20 Table 1 – Initial nitric acid reaction orates for toluene nitration in a 150 μm PTFE reactor. (Experiments used 78% H₂SO₄ with 7% HNO₃)

Figure 12 shows nitrotoluene production for a range of flow ratios and reactor lengths. The results show an increasing production of nitrotoluene when larger ratios of acid to organic were used in the reactor. However, the little improvement in conversion is observed for flow ratios exceeding 5:1 even though more acid is

available for nitration. This is also reflected in the lower reaction rates shown in Table 1 for the higher flow ratios. This would suggest a poorer mixing environment for the high flow ratios probably due to increased acid slug length.

- 5 End effects from possible post reactor nitration were also examined for both benzene and toluene nitration. Output from reactor tubes of different lengths were compared to check that increased length provided higher conversion implying that the reaction was taking place within the capillary tube and not within the sampling system. Figure 12 shows the results from three different reactor lengths using the same 10 conditions and shows in general that higher conversion was achieved for the longer tubes.

Visual analysis of liquid-liquid flow through a capillary reactor has shown that a pattern of alternating organic/aqueous slugs can be produced each having lengths 15 down to 300 μm . The work has also shown the importance of low internal volumes in distributor design for controlling the pattern of liquid-liquid flow produced.

Reaction results for benzene and toluene nitration have indicated reaction rates in the range of 1 to 8 min^{-1} can be produced from a capillary reactor. This would indicate 20 residence times for complete conversion to be in the region of 10 to 60 seconds. A comparison with some existing benzene nitration processes (as described in the indicated US patents) is shown in Table 2. This illustrates that even with 178 μm bore tubing the capillary reactor process is competitive.

Information source	Inlet (°C)	Outlet (°C)	H ₂ SO ₄ (mass %)	Conversion (%)	By-product (ppm)	Time (s)	Rate (min ⁻¹)
US 4,091,042	80	128	60.6	89.5	1000	120	0.9
US 4,091,042	80	134	65.2	99.1	2090	120	2.1
US 5,313,009	95	120	69.5	90	1750	25	4.6
Capillary 178µm	90	90	77.7	94.0	4600	24.4	5.90
Capillary 178µm	90	90	72.2	60.7	Below 1000	26.1	1.6

Table 2 – Comparison of benzene nitration performance
with existing processes

5

Ultimately, narrow channel microreactors based on this technique of liquid-liquid contacting will require shorter path lengths for diffusion to improve efficiency and lower by-product production. The use of microfabricated devices with more sophisticated distribution will be required to chop the liquids into smaller slugs or 10 droplets. Scale-up of the devices for chemical production will be achieved through use of parallel channels whilst their use for analysis will be facilitated through small on-chip versions.

Referring now to Figure 13, there is shown an alternative embodiment of the present 15 invention comprising two laminar plates 40, 41 which are mountable one 40 on top of the other 41 such that the plates 40, 41 are in registration with each other. The plates 40, 41 are made out of a non-stick material such as PTFE, and an upper surface 42 of plate 40 is provided with etched channels defining capillary pathways 43, 44, 45 which meet at a junction 46. Input capillary pathways 43, 44 meet substantially 20 head-on at the junction 46, and output capillary pathway 45 leads away therefrom substantially at right angles to the capillary pathways 43, 44. The other plate 41, when mounted on top of plate 40, provides a top surface for the capillary pathways

43, 44, 45. The plate 41 may be secured to the plate 40 by way of welding, adhesives, mechanical clamps or other suitable means.

5 Example 3: Acetic acid titration

Mass transfer performance of a glass capillary reactor distribution device having capillary pathway channels 0.38mm wide and 0.38mm deep in a standard configuration, as shown for example in Figure 13, was examined using a titration reaction. Kerosene loaded with 0.65 moles per litre of acetic acid was used in conjunction with an aqueous solution of sodium hydroxide at various concentrations of NaOH and containing phenol red pH indicator. Equal flow rates of each liquid phase were fed through the glass device and the time taken to transfer different quantities of acetic acid, as indicated by colour change in the aqueous system, was measured at two different flow velocities. Typical slug lengths produced were 1.1mm to 1.6mm long. The results are shown in Figure 14. These show a significant enhancement in performance gained by vortex mixing inside the slugs, compared with that expected from pure diffusion at this scale, and also that higher flow velocity produces faster mass transfer.

20

Finally, Figure 15 illustrates the flow dynamics of a pair of slugs 16, 17 passing along a capillary pathway 45 of a device as shown in Figure 14. The general flow direction is indicated by arrow A, with the internal circulation patterns being shown for each slug 16, 17, these patterns being caused by friction between the walls of the pathway 45 and boundary layers of the slugs 16, 17 as they progress along the pathway 45. Inter-slug diffusion occurs at an interface 50 between the slugs 16, 17.

Nomenclature

C	Reaction rate constant	s ⁻¹
D	Diffusivity	m ^{2.s⁻¹}
d	Path length for diffusion	m
Fo	Fourier number	-
t	Residence time	s
X	Proportion of nitric acid remaining	-

CLAIMS:

1. A capillary reactor distribution device comprising first and second capillary pathways which meet at a junction and a third capillary pathway which leads away from the junction, the capillary pathways being dimensioned such that, when first and second immiscible fluids are fed along respectively the first and second capillary pathways under predetermined laminar flow conditions, the first and second fluids chop each other into discrete slugs which pass along the third capillary pathway.
- 10 2. A device as claimed in claim 1, wherein the first and second capillary pathways meet substantially head on.
3. A device as claimed in claim 1, wherein the first and second capillary pathways meet substantially at right angles.
- 15 4. A device as claimed in claim 1, wherein the first and second capillary pathways meet at an angle between 90 and 180 degrees.
5. A device as claimed in claim 1, wherein the first and second capillary pathways meet at an angle between 0 and 90 degrees.
- 20 6. A device as claimed in claim 1, wherein the first and second capillary pathways meet at an angle from 90 to 300 degrees.
- 25 7. A device as claimed in any preceding claim, comprising a solid block into which the capillary pathways have been bored or otherwise formed.
8. A device as claimed in claim 7, wherein the solid block is made out of a non-stick material.
- 30 9. A device as claimed in claim 7 or 8, wherein the solid block is made out of a

material having a low surface energy.

10. A device as claimed in claim 7, wherein the capillary pathways are lined with a non-stick material.

5

11. A device as claimed in claim 7 or 10, wherein the capillary pathways are lined with a material having a low surface energy.

12. A device as claimed in any one of claims 7 to 11, wherein the capillary pathways extend from an interior portion of the solid block towards an outer surface thereof, and wherein attachment means are provided for attaching external capillary tubes to the capillary pathways.

13. A device as claimed in claim 12, wherein the attachment means are located on an outer surface of the solid body.

14. A device as claimed in claim 12, wherein the attachment means are located or extend within the solid block.

20 15. A device as claimed in any one of claims 12 to 14, wherein the attachment means include O-ring seals.

16. A device as claimed in claim 14 or 15, wherein the capillary pathways are defined by capillary tubes inserted into boreholes provided in the solid block and positioned so that mutually abutting ends of the capillary tubes form the junction.

25 17. A device as claimed in any one of claims 1 to 5, comprising at least two generally laminar plates mounted one directly on top of the other such that a surface of one plate contacts a surface of the other plate, at least one of the surfaces being provided with features serving to define the capillary pathways.

18. A device as claimed in claim 17, wherein the surface features comprise channels.

19. A device as claimed in claim 17 or 18, wherein the surface features comprise
5 ridge-like protrusions.

20. A device as claimed in any one of claims 17 to 19, wherein the plates are made of a non-stick material.

10 21. A device as claimed in any one of claims 17 to 20, wherein the plates are made of a material having a low surface energy.

22. A device as claimed in any one of claims 17 to 19, wherein the contacting surfaces of the plates are coated with a non-stick material.

15 23. A device as claimed in any one of claims 17 to 19 and 22, wherein the contacting surfaces of the plates are coated with a material having a low surface energy.

20 24. A device as claimed in any preceding claim, wherein at least one of the first and the second capillary pathways is provided with a filter.

25. A device as claimed in any preceding claim, wherein at least one of the first, second or third capillary pathways is lined with a chemically reactive material.

25 26. A device as claimed in claim 25, wherein the material is a solid heterogeneous catalyst.

27. A method for contacting two immiscible fluids, wherein a first fluid is fed
30 under laminar flow conditions along a first capillary pathway and a second fluid is fed under laminar flow conditions along a second capillary pathway, the first and

second capillary pathways meeting at a junction having a third capillary pathway leading away therefrom, and wherein the flow conditions in each of the first and second capillary pathways are selected such that the first and second fluids chop each other into discrete slugs which pass along the third capillary pathway.

5

28. A method according to claim 27, wherein the first and second capillary pathways meet substantially head on.

10 29. A method according to claim 27, wherein the first and second capillary pathways meet substantially at right angles.

30. A method according to claim 27, wherein the first and second capillary pathways meet at an angle between 90 and 180 degrees.

15 31. A method according to claim 27, wherein the first and second capillary pathways meet at an angle between 0 and 90 degrees.

32. A method according to claim 27, wherein the first and second capillary pathways meet at an angle from 90 to 300 degrees.

20

33. A method according to any one of claims 27 to 32, wherein the fluids are both liquids.

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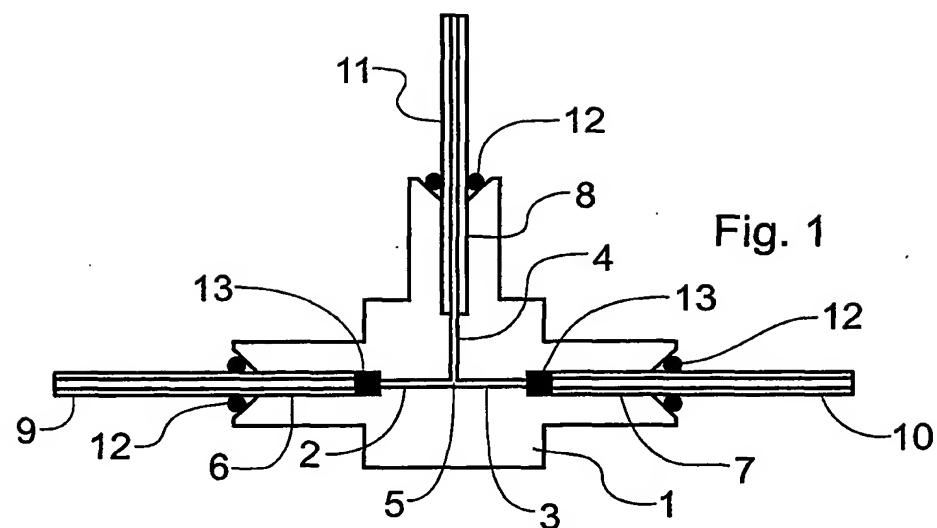


Fig. 1

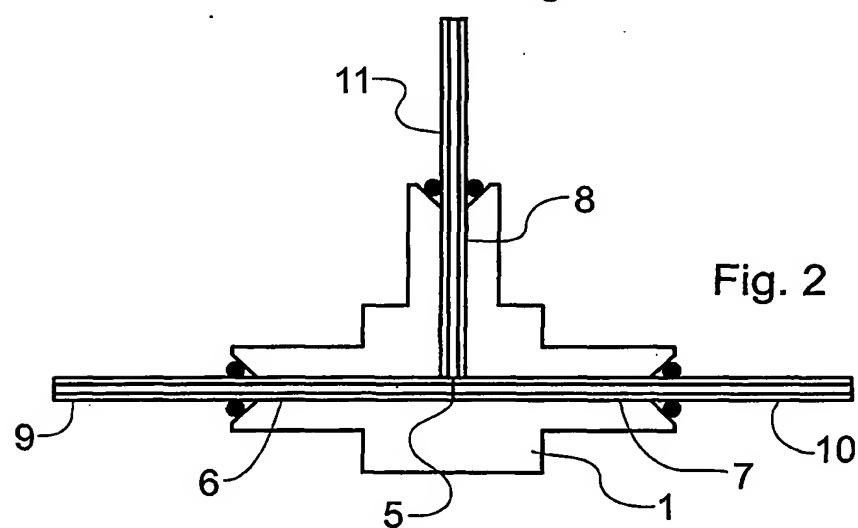


Fig. 2

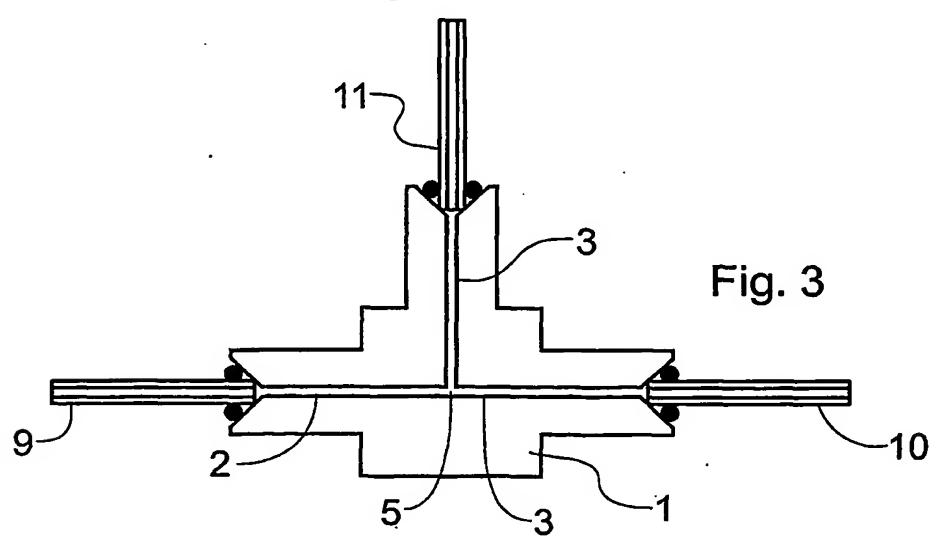


Fig. 3

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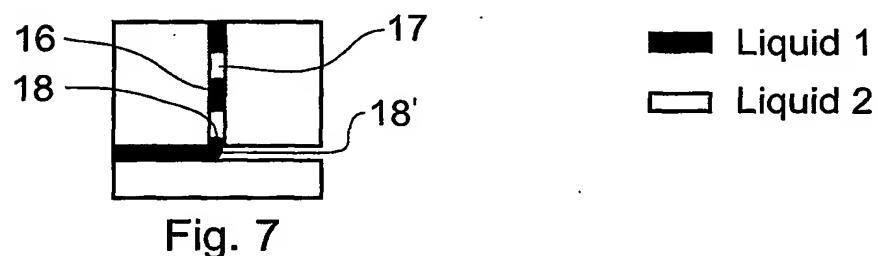
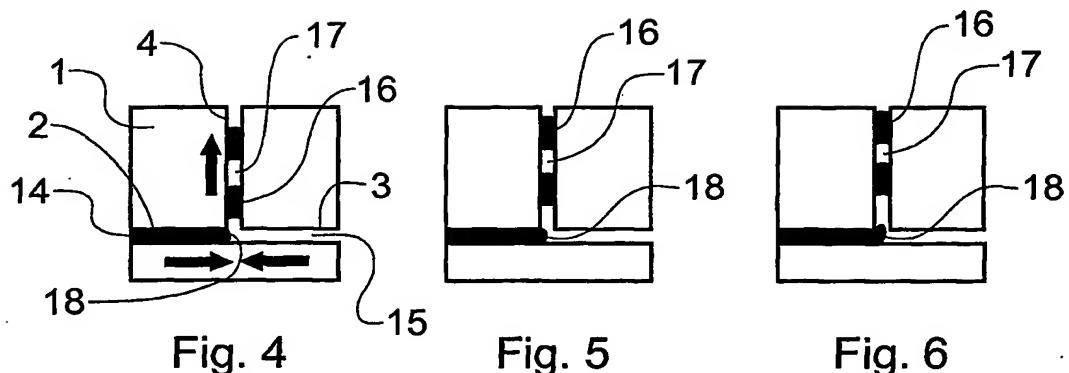


Fig. 8

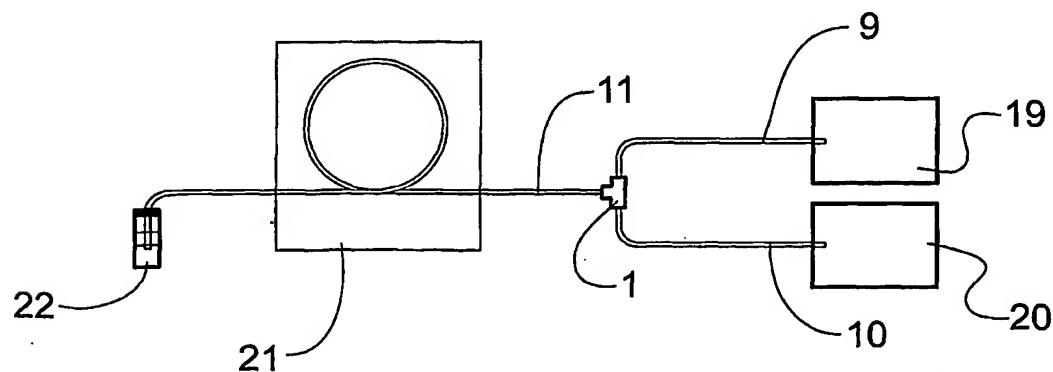


Fig. 9

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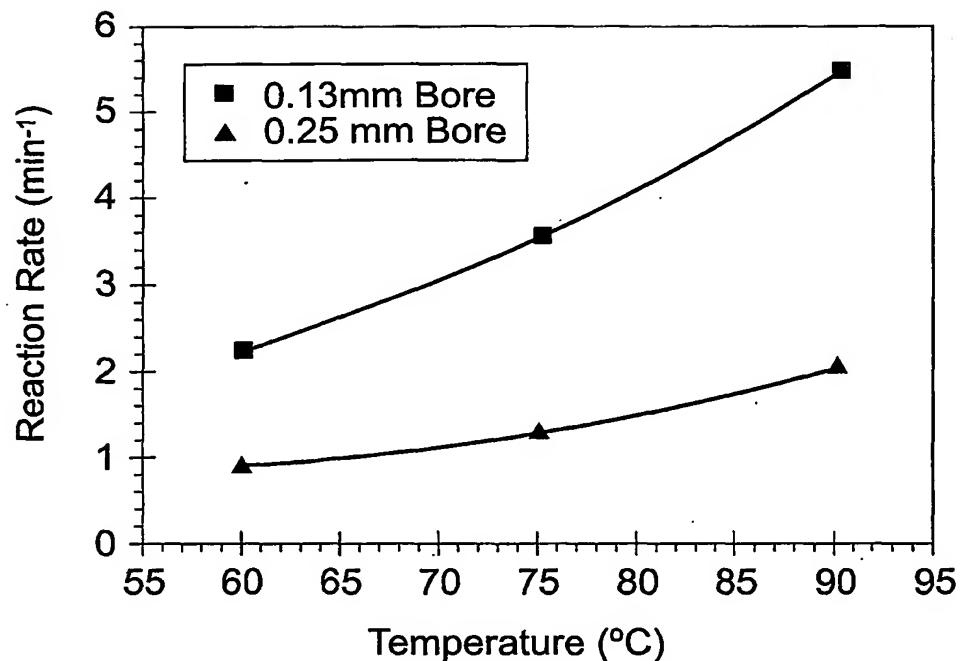


Fig. 10

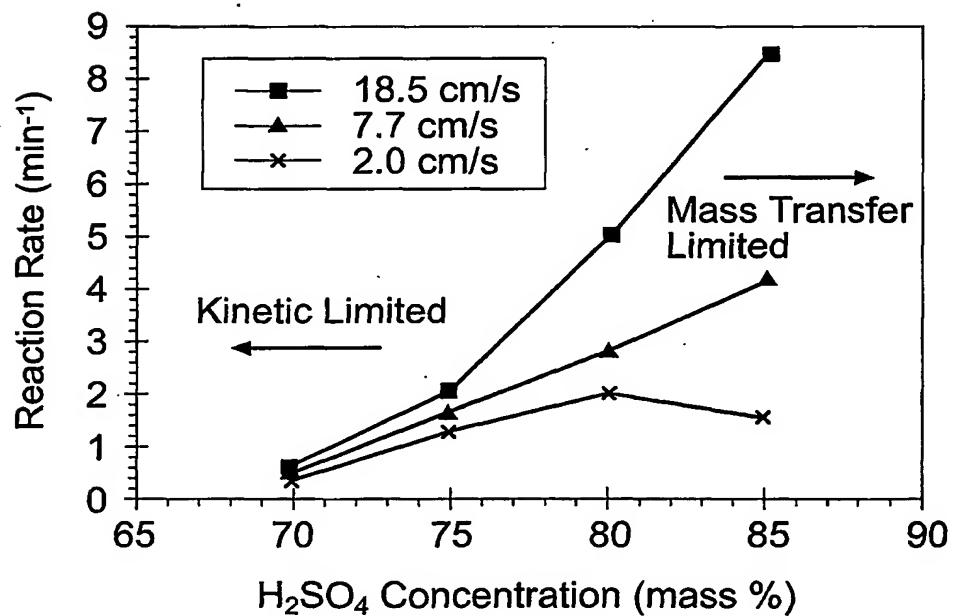


Fig. 11

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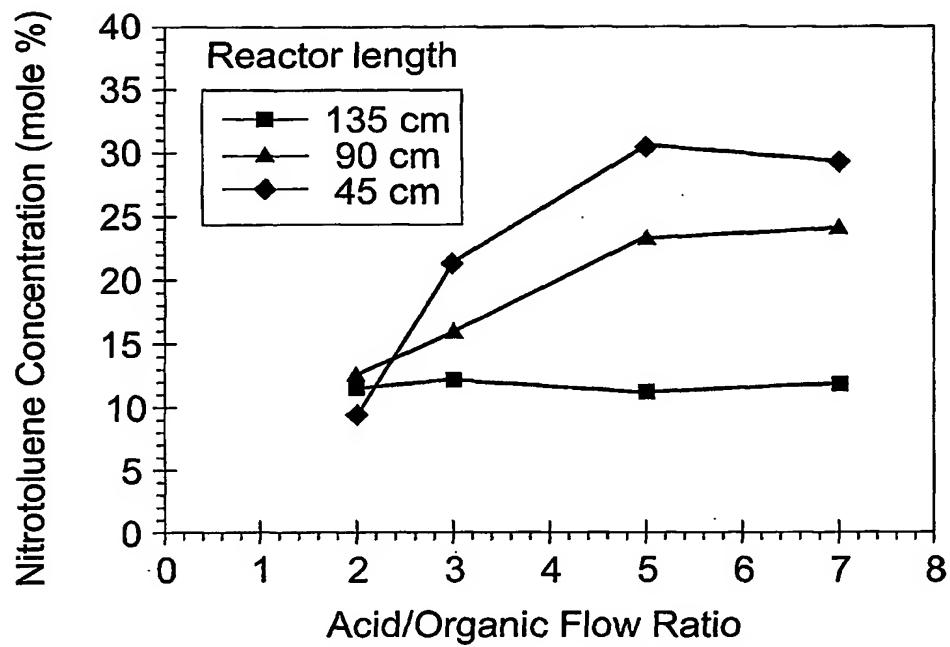


Fig. 12

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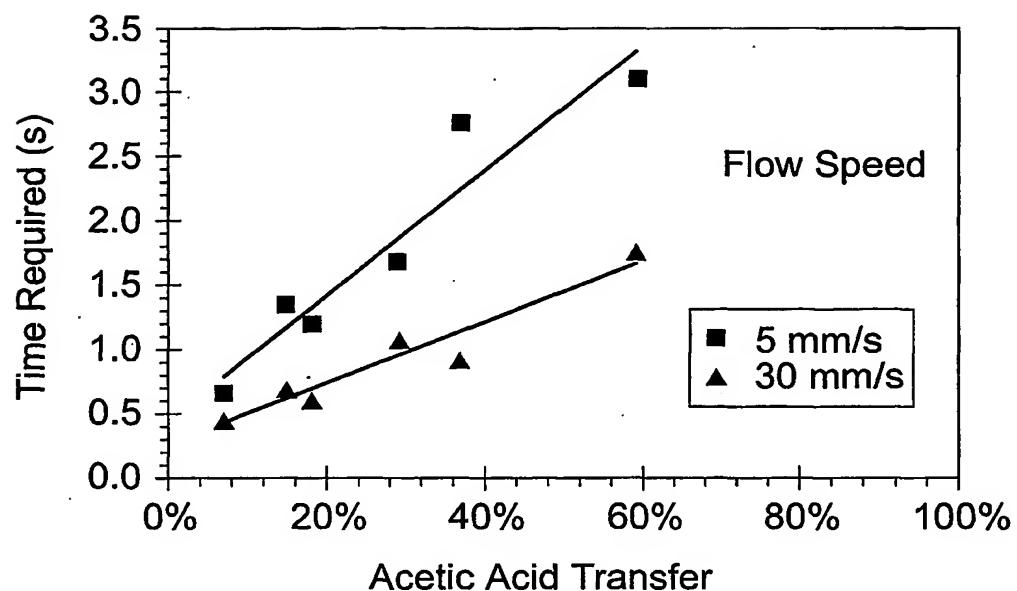


Fig. 13

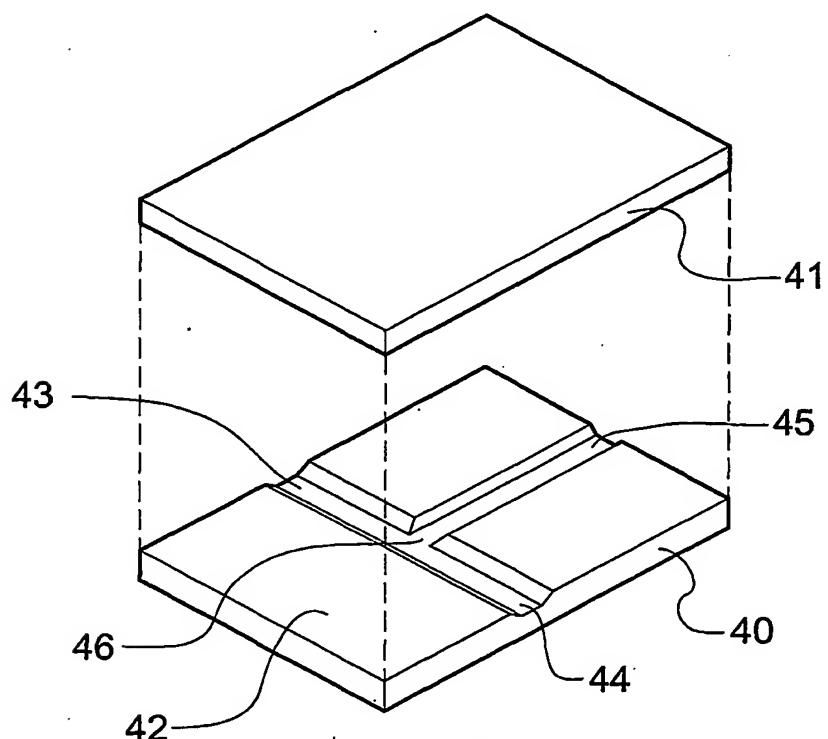


Fig. 14

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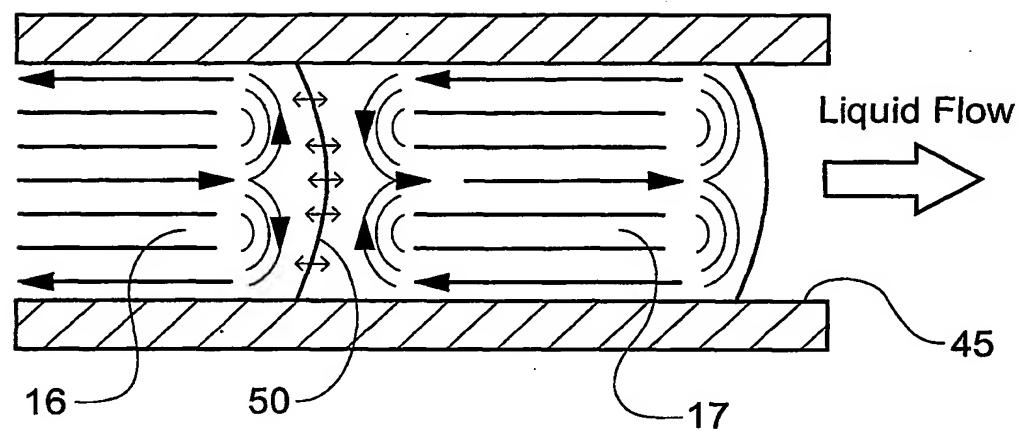


Fig. 15

INTERNATIONAL SEARCH REPORT

Interr	I Application No
PCT/GB 01/00848	

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7	B01J19/00	B01F5/02	B01F13/00	G01N35/08	B01L3/00
	F15C1/14				

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 B01J B01F G01N B01L F15C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 00442 A (THE UNIVERSITY OF WASHINGTON) 3 January 1997 (1997-01-03)	1,2,6,7, 12,13 27,28, 32,33
Y	page 5, line 28 -page 6, line 13 page 10, line 13 -page 11, line 17 page 12, line 27 -page 14, line 14 page 24, line 20 - line 27 figure 1 ----	
P,X	WO 00 61275 A (PENTH BERND) 19 October 2000 (2000-10-19) page 9, line 1 -page 13, line 2 figures 1-6 ----	1,2,4,6, 7,12,14, 16-18
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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Date of the actual completion of the International search	Date of mailing of the International search report
3 July 2001	11/07/2001
Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel: (+31-70) 340-2040, Tx: 31 651 epo nl. Fax: (+31-70) 340-3016	Vlassis, M

INTERNATIONAL SEARCH REPORT

Internat.	Application No
PCT/GB 01/00848	

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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A	column 2, line 25 -column 3, line 54 figure 2 ---	27,28, 32,33
Y	DE 195 36 103 A (DANFOSS AS) 3 April 1997 (1997-04-03) the whole document ---	27,28, 32,33
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Intern: Application No

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